

ARMICARE HAND SANITIZER- alcohol liquid

Armis Biopharma, Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ArmiCare Hand Sanitizer

DRUG FACTS

ACTIVE INGREDIENT

Ethyl Alcohol 75% v/v.

PURPOSE

Antiseptic

USES

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

WARNINGS

For external use only. Flammable. Keep away from heat or flame.

DO NOT USE

- On children less than 2 months of age
- On open skin wounds

When using this product

keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if

irritation or rash occurs. These may be signs of a serious condition.

KEEP OUT OF REACH OF CHILDREN

If swallowed, get medical help or contact a Poison Control Center right away at 1-800-222-1222

Directions:

- Place enough product on hands to cover all surfaces. Rub hands together until dry.

- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information:

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients:

Deionized water, Glycerin, Hydrogen Peroxide, Acetic Acid, Lavender, Polysorbate-20

QUESTIONS?

1-800-970-1779

Package Labeling:



ArmiCare™

Hand Sanitizer

Leaves Hands Feeling Silky Smooth

Contains 75% Ethyl Alcohol


Topical Solution

Net contents:
2 fl. oz. (60 mL)

ARMIS
BIOPHARMA™

ARMIS BIOPHARMA, Inc.
2950 E. Harmony Road, Suite 252
Ft. Collins, CO 80528
Tel: 970-266-5127

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DRUG FACTS (cont.)

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ARMICARE HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81793-187
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
ACETIC ACID (UNII: Q40Q9N063P)	
LAVANDULA ANGUSTIFOLIA SUBSP. ANGUSTIFOLIA FLOWER (UNII: 19AH1RAF4M)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:81793-187-00	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/05/2021	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph not final	part333E		04/05/2021	

Labeler - Armis Biopharma, Inc (166990189)

Revised: 4/2021

Armis Biopharma, Inc